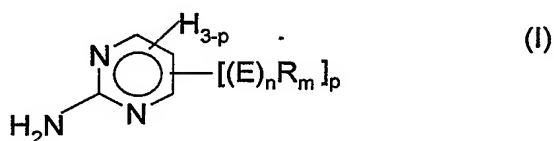
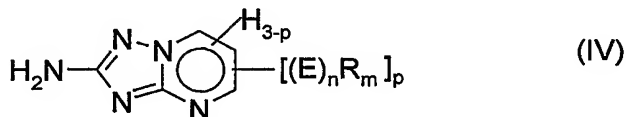


## Claims

1. Process for the preparation of unsubstituted or substituted 2-amino-[1,2,4]triazolopyrimidines which comprises combining A) 2-Amino-pyrimidine or its derivatives with alkyloxycarbonyl isothiocyanate or aryloxycarbonyl isothiocyanate with B) hydroxyl ammonium salt and a base wherein the reaction is carried out in a polar aprotic organic solvent in the temperature range of from 40 to 150 °C.
2. The process according to claim 1 wherein the pH value in step B) is increased over time and finally maintained in the range of from 5.5 to 7.5.
3. The process as in claims 1 to 2, wherein the hydroxylammonium salt is hydroxylammonium sulfate.
4. The process as in claims 1 to 3, wherein the polar aprotic solvent is selected from the group consisting of carboxylic acid esters.
5. The process as claimed in claims 1 to 4 wherein the 2-amino-pyrimidine or its derivatives is described by formula I.



and the 2-amino-[1,2,4]triazolopyrimidine is described by formula IV



wherein the variables have the following meaning:

E = independently the same or different are O, S, N, P;

R = independently the same or different are C<sub>1-10</sub>-alkyl; C<sub>6-20</sub>-aryl; C<sub>7-20</sub>-arylalkyl;

C<sub>7-20</sub>-alkylaryl which each of those may be substituted with one or more of the

following groups: F, Cl, Br, I, C<sub>1-20</sub>-alkoxy, C<sub>6-20</sub>-aryloxy, non substituted or preferably substituted amino; F, Cl, Br, I;

- n = 0 or 1  
m = 1 for E = O, S  
m = 2 for E = N, P  
5 p = 0, 1, 2 or 3.
6. Process as claimed in claims 1 to 5, wherein the process is conducted without isolation of intermediates.
- 10 7. Process for the preparation of N-([1,2,4]triazolo[1,5-a]pyrimidin-yl)aryl sulfonamides or N-([1,2,4]triazolo[1,5-a]pyrimidin-yl)heteroaryl sulfonamides which comprises preparing unsubstituted or substituted 2-amino-[1,2,4]triazolopyrimidines according to claim 1 to 6 and subsequently reacting the yielded unsubstituted or substituted 2-amino-[1,2,4]triazolopyrimidines with an arylsulfonylhalogenide Ar-  
15 SO<sub>2</sub>-Hal or an heteroarylsulfonylhalogenide Hetar-SO<sub>2</sub>-Hal.
8. Use of a process as claimed in claims 1 to 6 in the synthesis of N-([1,2,4]triazolo[1,5-a]pyrimidin-yl) structure containing agrochemicals or pharmaceuticals.

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